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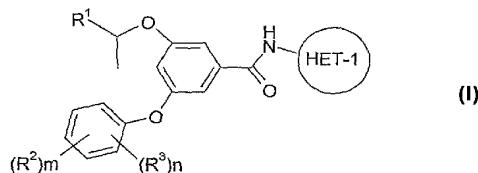
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(54) Title: BENZAMIDE DERIVATIVES AND THEIR USE AS GLUCOKINASE ACTIVATING AGENTS



(57) Abstract: Compounds of Formula (I): wherein: R<sup>1</sup> is methoxymethyl; R<sup>2</sup> is selected from -C(O)NR<sup>4</sup>R<sup>5</sup>, -SO<sub>2</sub>NR<sup>4</sup>R<sup>5</sup>, -S(O)<sub>p</sub>R<sup>4</sup> and HET-2; HET-1 is a 5- or 6-membered, optionally substituted C-linked heteroaryl ring; HET-2 is a 4-, 5- or 6-membered, C- or N-linked optionally substituted heterocycl ring; R<sup>3</sup> is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano; R<sup>4</sup> is selected from for example hydrogen, optionally substituted (1-4C)alkyl and HET-2; R<sup>5</sup> is hydrogen or (1-4C)alkyl; or R<sup>4</sup> and R<sup>5</sup> together with the nitrogen atom to which they are attached may form a heterocycl ring system as defined by HET-3; HET-3 is for example an optionally substituted N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocycl ring; p is (independently at each occurrence) 0, 1 or 2; m is 0 or 1; n is 0, 1 or 2; provided that when m is 0, then n is 1 or 2; or a salt, pro-drug or solvate thereof, are described. Their use as GLK activators, pharmaceutical compositions containing them, and processes for their preparation are also described.

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